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• IKEDA, Shin
Katano-shi, Osaka 576-0022 (JP)
• YOSHIOKA, Toshihiko
Hirakata-shi, Osaka 573-0035 (JP)

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(74) Representative:
Schuster, Thomas, Dipl.-Phys.
Grünecker, Kinkeldey, Stockmair &
Schwanhäusser
Anwaltssozietät
Maximilianstrasse 58
80538 München (DE)

(71) Applicant:
Matsushita Electric Industrial Co., Ltd.
Kadoma-shi, Osaka 571-8501 (JP)

(72) Inventors:
• WATANABE, Motokazu
Kadoma-shi, Osaka 571-0064 (JP)

(54) BODY FLUID TESTING DEVICE

(57) A body fluid testing device capable of a high-accurate measurement by using a small amount of body fluid and obtained by additionally providing a pump for evacuating the inside of a device body to a negative pressure and a three-way solenoid valve for releasing the negative pressure or additionally providing a humidifying means for supplying steam into the device body to a body fluid testing device comprising a container-shaped testing device body having an opening, a sensor for testing a body fluid and a body fluid exuding device.

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Description**Technical Field**

[0001] The present invention relates to an examination system which oozes bodily fluids on skin surface and collects the oozed fluids, then examines the fluids with a sensor.

Background Art

[0002] When blood sugar is examined in a conventional manner, an oozing device for bodily fluids such as a lancet is used for pricking skin in order to have the bodily fluids ooze there. Then an examination device for bodily fluids is applied there to collect specimen. This process requires numbers of operations before the examination is completed, and thus an automated examination system which sequentially carries out these operations is demanded.

[0003] One of the automated examination systems is disclosed in the Japanese Patent Application Non-examined Publication No. H10-33508. This system includes a tightening belt into which a finger is inserted, a needle for pricking the finger tip, and a sensor disposed nearby the needle. After being pricked with the needle, the finger is tightened with the belt so that bodily fluids ooze, and contact with the sensor.

[0004] However, when the bodily fluids ooze over the skin surface, the fluids sometimes does not reach to the sensor, and this system thus cannot carry out the examination.

[0005] Further, because the bodily fluids are vaporized to a degree before they reach to the sensor, a precise examination is not expected particularly when the bodily fluids oozed are in a little quantity.

[0006] The present invention addresses the problems discussed above, and aims to provide an examination system for bodily fluids. This system of the present invention can supply the bodily fluids free from errors to a specimen collector of a sensor, even when the bodily fluids oozed by the examination system are in a little quantity.

[0007] Another object of the present invention is to provide an examination system for bodily fluids, and which suppresses evaporation from the bodily fluids oozed so that the examination can produce a precise result.

Summary of the Invention

[0008] An examination system of the present invention aims to solve the problems discussed above and comprises the following elements:

a vessel-like examination unit having an opening in front;
decompressing means for decompressing the

inside of the unit; and

a specimen collector provided in the unit and facing the opening.

5 [0009] The present invention also provides an examination system which includes a bodily-fluids-oozing-device in the unit discussed above.

[0010] The present invention provides another examination system which comprises the following elements:

a vessel-like examination unit having an opening in front;

humidifying means for humidifying the inside of the unit;

15 a bodily-fluids-oozing-device; and

a specimen collector provided in the unit and facing the opening.

[0011] The present invention further provides an examination system which includes decompressing means facing the opening in the examination unit.

[0012] The specimen collector preferably incorporates with a sensor, and a specimen-supplying-outlet of the device in the unit preferably faces the opening.

[0013] The present invention still further provides an examination system which comprises the following elements:

a vessel-like examination unit having an opening in front;

30 decompressing means for decompresses the unit; decompress releasing means for restoring a decompressed unit to atmospheric pressure; a bodily-fluids-oozing-device;

35 a specimen collector provided in the unit and facing the opening;

a sensor having an electrode system for contacting with the specimen supplied from the specimen collector and outputting information about the specimen as an electric signal from the electrode system;

40 determining means for determining a measured value with the electric signal; and

45 a controller for controlling the decompressing means, decompress-releasing-means and sensor.

This examination system allows the decompress-releasing-means to function after the sensor outputs an electric signal. It is preferable for this system that the decompressing means functions after the bodily-fluids-oozing-device functions as well as the decompress-releasing-means functions after the sensor outputs an electric signal.

[0014] The present invention yet provides another examination system which comprises the following elements:

a vessel-like unit having an opening in front;

humidifying means for humidifying the inside of the unit;
 decompressing means for decompressing the inside of the unit;
 decompress releasing means for restoring a decompressed condition to atmospheric pressure; a bodily-fluids-oozing-device;
 a specimen collector provided in the unit and facing the opening;
 a sensor having an electrode system for contacting with the specimen supplied from the specimen collector and outputting information about the specimen as an electric signal from the electrode system;
 determining means for determining a measured value with the electric signal; and
 a controller for controlling the humidifying means, decompressing means, decompress-releasing-means and sensor.

This examination system allows the bodily-fluids-oozing-device to function after the humidifying means functions as well as the decompress-releasing-means to function after the oozing device functions, and the decompress-releasing-means to function after the sensor outputs an electric signal.

[0015] The examination systems discussed above produce the following advantages:

[0016] In one of the embodiments of the systems, urge the opening of unit against skin—from which bodily fluids are to be sucked—thereby blocking up the opening, then have the decompressing means decompress the inside of the unit for rising the skin within the opening so that the bodily fluids can be supplied to the specimen collector with ease.

[0017] The bodily-fluids-oozing-device, which oozes bodily fluids from the skin, can be arranged in the examination unit so that the opening of the unit faces the oozing device, whereby a series of operations such as oozing and sucking bodily fluids can be sequentially carried out.

[0018] In another embodiment of the systems, urge the opening of unit against skin—from which bodily fluids are to be sucked—thereby blocking up the opening, then have the humidifying means humidify the inside of the unit. This prevents the bodily fluids oozed by the bodily-fluids-oozing-device from being vaporized. The oozing device is disposed in the unit so that the evaporation can be effectively suppressed.

[0019] When the inside of the unit is decompressed, the bodily fluids are subject to evaporation, this system thus allows the humidifying means to work before the decompressing means starts to work in order to humidify the inside of the unit in advance.

[0020] As the humidifying means, e.g. a water bowl—having a heater or an ultrasonic-wave-generator on the bottom—can be used. The upper section of the bowl communicates with the examination unit so that

vapor produced from the water in the bowl flows into the unit, in particular, into the opening side.

[0021] A fan disposed at the communicating section between the unit and the bowl can efficiently run the vapor produced in the bowl into the unit.

[0022] When a humidity inside the unit rises so high, the inner wall of the unit and a surface of the sensor holder are sometimes dewed. If the dews drop onto the skin surface, the bodily fluids are thinned. This adversely affects the examination result. It is therefore preferable to dispose a humidity-sensor-probe in the unit for monitoring the humidity therein when bodily fluids are examined. Considering an effective prevention against the evaporation, the humidity in the unit preferably ranges from 60 to 70%.

[0023] When a decompressing pump is used as the decompressing means, the inside of the unit is quickly decompressed so that the bodily fluids can be quickly supplied to the specimen collector.

[0024] Further, when decompress-releasing-means is disposed in the unit for restoring the decompressed condition inside the unit to atmospheric pressure, it prevents air from flowing rapidly into the unit at removing the unit from the skin after the examination. As a result, the bodily fluids are not scattered.

[0025] An operation program of the examination system can be set so that the decompress-releasing-means works at the same time on or with some delay of completing the examination. This can notify a user of weak eyes with a feeling of skin stretch that the user is still under the examination.

Brief Description of the Drawings

[0026]

Fig. 1 is a cross section illustrating a schematic construction of an examination system for bodily fluids in accordance with a first exemplary embodiment of the present invention.

Fig. 2 is a cross section of an essential part of a decompress-releasing device of the examination system shown in Fig. 1.

Fig. 3 is a block diagram illustrating a circuit of the examination system in accordance with a second exemplary embodiment of the present invention.

Fig. 4 is a flowchart illustrating an operation of the examination system.

Fig. 5 is a cross section illustrating part of schematic construction of an examination system in accordance with a third exemplary embodiment of the present invention.

Fig. 6 is a block diagram illustrating a circuit of an examination system in accordance with a fourth exemplary embodiment of the present invention.

Fig. 7 is a flowchart illustrating an operation of the examination system.

Description of the Preferred Embodiments

[0027] Exemplary embodiments of the present invention are demonstrated hereinafter with reference to the accompanying drawings.

(Exemplary Embodiment 1)

[0028] Fig. 1 shows a construction of an examination system for bodily fluids in accordance with the first exemplary embodiment of the present invention.

[0029] Examination unit 1 comprises cylinder 2 having a bottom plate and cap 3 screwed detachably on the opening of cylinder 2. An opening of the unit, formed at the tip of cap 3, is e.g. an oval having 15 mm longer diameter and 10 mm shorter diameter. This opening is equipped with adapter 4 made of rubber to protect skin and achieve tighter adherence to the skin.

[0030] Sensor holder 5 and bodily-fluids-oozing-device 6 are disposed inside of the unit 1. Sensor holder 5 is mounted with sensor 19 at its tip, and includes a terminal contacting with an electrode terminal of the sensor and a lead-wire coupling this terminal to an examining section (not shown). Oozing device 6 comprises needle holder 8 to which needle supporter 7 is mounted, while a needle is fixed to the tip of needle supporter 7, tube 9 guiding needle holder 8, spring 10 protruding needle holder 8, and switch 11.

[0031] Needle holder 8 is unitarily formed with stopper 12 which is urged toward left in the drawing by stopper's own elasticity. This stopper 12 engages with step 13 provided in tube 9 so that needle holder 8 remains within tube 9. When switch 11 presses stopper 12 rightward and releases the engagement with step 13, needle holder 8 protrudes to the opening of the unit by the elasticity of spring 10.

[0032] A closed top section of unit 1 is coupled to electromagnetic cross-valve 15 by pipe 14. Electromagnetic valve 15 is coupled to decompressing pump 18 via pipe 17 and to releasing pipe 16. In this embodiment, sensor holder 5 is equipped with sensor 19 for measuring blood sugar. A specimen-supplying-outlet at the tip of sensor 19 works as a specimen collector. The specimen supplying outlet sucks blood by capillarity.

[0033] A tip of oozing device 6 is away from skin 20 comparing with the specimen collector of sensor 19. The bodily fluids ooze have thus no chance to contact with the tip of device 6 when the inside of unit 1 is decompressed. As shown in Fig. 1, it is preferable to set a place—to be pricked by the needle of oozing device 6—at or near the center of the opening of the unit because bodily fluids 21 are accumulated in a top of skin swelling up as marked with 60. It is also preferable to arrange the specimen collector of the sensor near to the center of the unit's opening because a space between the collector and the skin becomes shorter.

[0034] The opening of the unit is preferably shaped like a round or an oval rather than a square in order to

block up the opening with the skin airtightly. The unit per se is preferably made of transparent material such as glass or plastic, and plastic is more preferable from the standpoint of light weight and safety. As the decompressing means, an electric-driven decompressing pump is preferable because of its strong decompressing power. A vacuum pump is more preferable among others because of its smaller size.

[0035] An operation and an effect of this examination system are demonstrated hereinafter.

[0036] First, press the opening of unit 1 against skin 20 such as a finger thereby blocking up the opening. Next, prick the skin with oozing device 6 so that bodily fluids are oozed and sucked. In this embodiment, blood is taken as an example and is sucked. The needle supported by needle supporter 7 is biased by spring 10 and reaches to the skin, then pricks slightly the skin surface. Blood thus oozes from the skin surface. At this moment, decompressing pump 18 coupled to the unit lowers the inside pressure of the unit from atmospheric pressure. The skin swells up toward the inside of the unit due to this decompress, and the blood on the skin surface contacts with the specimen collector of the sensor, then the blood is supplied to the sensor. The finger tip swells to a degree; however, the decompressing means raises the skin so that the blood can contact with the specimen collector of the sensor, thus individual differences in height of the skin raised does not affect the specimen collection. Further, when a user of diabetes, e.g. and being suffered from the weak eye due to the complications, uses this examination system, the bodily fluids are supplied to the sensor free from failures. Because examination unit 1 can be moved arbitrarily, the user can prick any spot of the body such as finger, palm, back of the hand, arm, belly, earlobe and the like.

[0037] In the first exemplary embodiment, bodily-fluids-oozing-device 6 uses the needle biased by spring 10; however, laser beam and the like, which prick slightly the skin surface, can be used with the same effect.

[0038] The sensor is disposed within the unit; however, as far as the specimen collector is disposed within the unit, the sensor per se can be disposed outside of the unit.

[0039] The bodily-fluids-oozing-device is not always equipped to unit 1. When the device is not equipped to the unit, an oozing device available on the market, e.g. a lancet, is used to ooze the bodily fluids. Then the unit is applied to the spot where the bodily fluids ooze, and the decompressing means decompresses the inside of the unit thereby raising the skin for supplying the bodily fluids to the sensor.

[0040] Electromagnetic cross-valve is used as the decompress-releasing-means in this first embodiment; however, the decompress-releasing-means separated from the cross valve can be provided to the unit. As shown in Fig. 2, for instance, pipe 22 communicating with the inside of the unit is provided to a top section of

unit 1, and rotary valve 23 is mounted to this pipe. Rotary valve 23 is rotated manually to have air-hole 25 communicate with air-permeable-hole 24 punched on pipe 22 so that the decompressed status in examination unit 1 can be released.

[0041] If unit 1 is removed from the skin being kept the decompressed status, the air flows into unit 1 abruptly thereby scattering the bodily fluids oozed. This may dirty the circumference. The decompress-releasing-means thus releases the decompressed condition so that the pressure inside the unit is restored to atmospheric pressure, and the bodily fluids have no chance to scatter. Further, this arrangement allows a user to remove the unit from the skin with ease free from a forcible manner. When the skin is stretched as strong as a user feels a pain due to decompressing, the rotary valve is rotated thereby releasing quickly the decompressed condition to remove the unit from the skin with ease.

[0042] The bodily fluids—which this examination system examines—include blood, lymphatic fluid, intercellular fluid and perspiration, which can be collected from skin surface.

(Exemplary Embodiment 2)

[0043] Fig. 3 is a block diagram illustrating a circuit of an examination system in accordance with the second exemplary embodiment.

[0044] A diabetic patient measures his own blood sugar by himself as a routine work. The patient is sometimes suffered from detachment of retina, one of complications of diabetes, and in the worst case the patient loses his eyesight. Even if the patient eventually has weak eyes, he must measure the blood sugar routinely. When a patient of weak eyes examines his own blood, it is not easy for the patient to determine whether the blood oozes on the skin or not.

[0045] According to the second embodiment, a patient of weak eyes can examine his own blood free from failures using a sensor with an auto-start function and decompressing means for decompressing the inside of the unit. In this second embodiment, a needle is used as bodily-fluids-oozing-device same as the first embodiment and a vacuum pump is used as the decompressing means. Blood is taken as an example of bodily fluids to be examined. The vacuum pump is coupled to the unit as shown in Fig. 1 and can decompress the inside of the unit.

[0046] In Fig. 3, examination unit 1, to which a sensor is mounted for collecting and examining the bodily fluids, communicates with electromagnetic valve 55 and vacuum pump 56 via pipes. Computer 51 controls electromagnetic valve 55 and vacuum pump 56 thereby decompressing the inside of the unit or releasing the decompressed condition. Computer 51 further applies a voltage to electrode system 57 provided to the sensor and receives an electric signal from electrode system 57. It also issues a next instruction based on the electric

signal, determines a measured value by calculating the electric signals, and outputs the instruction as well as the measured value on display 54.

[0047] Computer 51 is initialized by turning main-switch 52 on and is loaded with a program which drives vacuum pump 56 by activating the bodily-fluids-oozing-device.

[0048] An operation of this system is demonstrated with reference to the flowchart shown in Fig. 4.

[0049] First, apply the opening of unit 1 to the skin, and turn on main switch 52 of computer 51 to initialize the system, such as clearing the memories (Step 101). Then turn on switch 53 of the bodily-fluids-oozing-device to prick the skin with the needle (Step 102). At this moment, if the needle substantially reaches to the skin, blood oozes.

[0050] After a while of switch 53 being turned on, vacuum pump 56 discharges the air from unit 1, i.e. decompress-operation starts (Step 103).

[0051] In step with the progress of decompressing operation inside of the unit, the skin urged tightly to unit 1 is sucked and swells up inside unit 1 and eventually contacts with the specimen collector of the sensor. At this moment, if the blood has oozed, the blood is supplied to the specimen collector. A voltage has been applied between at least two electrodes of the sensor, and computer 51 detects the availability of specimen through fluid communication between the electrodes when the blood is supplied to the sensor (Step 104), and starts measuring (Step 105). After the measurement, decompress-releasing-means releases the decompressed condition in unit 1 (Step 106). After the condition inside the unit is restored to atmospheric pressure, examination unit 1 is removed from the skin, and the examination is completed.

[0052] In Step 104, when computer 51 does not detect fluid communication between the electrodes and then determines that no blood oozes on the skin surface after a given period e.g. 15 seconds of starting the decompression, the decompressed condition in unit 1 is released (Step 107). Further, this is displayed on display 54, or alarm sound is produced so that an examiner can notice that the blood is not collected. Because the needle insufficiently pricks the finger, the system cannot collect the blood. Therefore, urge the finger against unit 1, for instance, to shorten the distance between the skin and the needle, and repeat Step 102 and onward.

[0053] When another sensor—which measures a specific component in the bodily fluids with the amount of current running between the electrodes of the electrode system—is used, an intense shock to the sensor would run noise current, which adversely affects the examination. Since a user of weak eyes, in particular, cannot monitor the progress of examination by his eyes, this adverse-influence sometime prevents the examination from being carried out in a normal way.

[0054] In this second embodiment, the inside of unit 1 is thus decompressed before the sensor starts to meas-

ure, and the decompressed condition is released after the measurement is completed. In other words, the examination system is programmed so that the decompress-releasing-means works at the same time or after some delay when the examination is completed. This arrangement allows a user to feel that he is still under the examination although he cannot monitor whether the needle pricks his skin or not in Step 102. Because the inside of unit 1 is decompressed before the sensor starts to measure, the user can feel that he is under the examination in Step 103, where the decompress operation starts, through Step 106, where the decompressed condition is released.

[0055] As a result, a user of weak eyes can avoid applying shocks to the examination system during the examination and prevent the adverse influence to the bodily-fluids examination.

[0056] When decompressing and releasing thereof are repeated often, a motor-driven pump is preferably used for the decompressing means because it starts decompressing promptly after power-on.

(Exemplary Embodiment 3)

[0057] Fig. 5 illustrates a structure of an examination system for bodily-fluids in accordance with the third exemplary embodiment.

[0058] Examination unit 31 comprises double-cylinder with a bottom plate, and cap 3 detachably screwed on an opening of inner cylinder 32. Outer cylinder 33 blocks up its lower end by itself. An opening of unit 31 is formed at tip of cap 3 and is equipped with adapter 4 made of rubber.

[0059] Outer cylinder 33 communicates with inner cylinder 32 via blower 38 and air hole 39. Blower 38 comprises pipe 41 and fan 40 disposed in pipe 41.

[0060] A humidifier comprises blower 38, outer cylinder 33, water bowl 34 disposed within cylinder 33, and heater 35 disposed within the bowl. Water bowl 34 is filled up with water 36. On the outer wall of cylinder 33, water-supply-outlet is provided for supplying water 36 into bowl 34. When the water is supplied, cap 37 is removed from the outlet.

[0061] Sensor holder 5, bodily-fluids-oozing device 43, and humidity-sensor probe 42 are disposed inside of cylinder 32.

[0062] Humidity-sensor probe 42 has a lead-wire coupling the probe to a controller (not shown). Sensor holder 5 is equipped with sensor 19 at its tip, and has a terminal contacting with an electrode terminal of the sensor as well as a lead-wire coupling the terminal to an examining section (not shown).

[0063] Bodily-fluids-oozing device 43 comprises a laser beam radiator which irradiates laser beam by depressing switch 44, and has a lead wire coupling the device 43 to a power supply (not shown).

[0064] Laser beam radiator 43 is placed so that laser beam hits the center or around the center of the opening

of unit 31. Sensor 19 is arranged to face the opening center so that a specimen supplier of the sensor is located near the opening-center of unit 31.

[0065] An operation and an effect of this examination system is demonstrated hereinafter.

[0066] First, warm up water 36 in bowl 34 by heater 35 to raise the humidity in the humidifier. Then blow the air of the humidifier into cylinder 32 by fan 40 to humidify the inside of cylinder 32. In this condition, urge the opening of unit 31 against skin 20, e.g. a finger, to block up the opening. Then control the humidity inside cylinder 32 at ca. 65% with humidity-sensor probe 42.

[0067] Next, depress switch 44 to irradiate laser beam, which pricks the skin surface and results in oozing bodily fluid 21. Then decompress the inside of cylinder 32 with decompressing pump 18 coupled to cylinder 32. The skin swells up toward the inside of the unit and the bodily fluid contacts with the specimen supplier. Finally the bodily fluid is supplied to the sensor.

(Exemplary Embodiment 4)

[0068] Fig. 6 is a block diagram illustrating a circuit of an examination system for bodily fluids in accordance with the fourth exemplary embodiment.

[0069] In Fig. 6, a sensor for collecting and examining the bodily fluids is mounted to examination unit 31, which communicates with electromagnetic valve 55 as well as vacuum pump 56 via pipes. Computer 51 controls valve 55 and pump 56 to decompress the inside of unit 31 or release the decompressed condition.

[0070] Computer 51 applies a voltage to electrode system 57 provided to the sensor and receives an electric signal from electrode system 57. It also issues a next instruction based on the electric signal, calculates the electric signals and then determines a measured value, and outputs the instruction as well as the measured value on display 54.

[0071] Computer 51 also receives the electric signal regarding the humidity from humidity-sensor probe 42, and based on the humidity information it controls on-off operation of the fan of blower 38 as well as on-off operation of heater 35.

[0072] Computer 51 is initialized when main-switch 52 is turned on, and then allows heater 35 and blower 38 to humidify the inside of unit 31. When the computer determines that the humidity is not less than 50% based on the humidity information from sensor probe 42, the computer is programmed to firstly activate the laser beam radiator and then drive vacuum pump 56.

[0073] For instance, when humidity-sensor probe 42 detects a humidity as high as 80%, computer 51 is programmed to turn off the fan of blower 38 and heater 35.

[0074] An operation of this system is demonstrated with reference to the flowchart shown in Fig. 7.

[0075] First, turn on main-switch 52 of computer 51 to initialize the system, which includes clearing the memories (Step 101).

[0076]	Second, power on heater 35 to warm up water in the bowl, then activate blower 38 to humidify the inside of unit 31 (Step 108). This humidifying is preferably started immediately after the system is initialized because the inside of unit 31 has been desirably humidified at the laser beam radiation in order to avoid vaporizing the bodily fluids.	10	Spring
		11	Switch
		12	Stopper
		13	Step
5		14	Pipe
		15	Electromagnetic cross-valve
		16	Releasing pipe
		17	Pipe
		18	Decompressing pump
10		19	Sensor
		20,60	Skin
		21	Bodily fluids
		22	Tubular section
		23	Rotary valve
15		24	Air-permeable-hole
		25	Air hole
		31	Entire system
		32	Inner cylinder
		33	Outer cylinder
20		34	Water bowl
		35	Heater
		36	Water
		37	Cap
		38	Blower
25		39	Air hole
		40	Fan
		41	Pipe
		42	Humidity-sensor-probe
		43	Laser beam radiator
30		44	Switch
		51	Computer
		52	Main switch
		53	Switch
		54	Display
35		55	Electromagnetic valve
		56	Vacuum pump
		57	Electrode system
	Claims		
40	1.		A bodily-fluids-examination-system comprising:
			a vessel-like examination unit having an opening in front;
45			decompressing means for decompressing an inside of said unit; and
			a specimen collector disposed in said unit and facing the opening.
50	2.		The bodily-fluids-examination-system as defined in Claim 1 further comprising a bodily-fluids-oozing-device.
55	3.		A bodily-fluids-examination-system comprising:
			a vessel-like examination unit having an opening in front;
			humidifying means for humidifying an inside of

- said unit;
a bodily-fluids-oozing-device; and
a specimen collector disposed in said unit and facing the opening.
4. The bodily-fluids-examination-system as defined in Claim 3 further comprising decompressing means for decompressing the inside of said unit. 5
5. The bodily-fluids-examination-system as defined in Claim 1, 2 or 4 further comprising decompress-releasing-means for restoring a decompressed condition in said unit to atmospheric pressure. 10
6. The bodily-fluids-examination-system as defined in Claim 2, 3 or 4 wherein said bodily-fluids-oozing-device includes one of a needle and a laser beam radiator. 15
7. The bodily-fluids-examination-system as defined in Claim 5 wherein said bodily-fluids-oozing-device includes one of a needle and a laser beam radiator. 20
8. A bodily-fluids-examination-system comprising:
a vessel-like examination unit having an opening in front;
decompressing means for decompressing an inside of said unit;
decompress-releasing-means for restoring a decompressed condition in said unit to atmospheric pressure; 30
a bodily-fluids-oozing-device;
a specimen collector disposed in said unit and facing the opening; 35
a sensor, having an electrode system for contacting with a specimen supplied from said specimen collector, for outputting information regarding the specimen as an electric signal from the electrode system;
determining means for determining a measured value of the specimen based on the electric signal; 40
a controller for controlling said decompressing means, said decompress-releasing-means and said sensor, wherein said system allows said decompress-releasing-means to function after said sensor outputs the electric signal. 45
9. A bodily-fluids-examination-system comprising:
a vessel-like examination unit having an opening in front;
decompressing means for decompressing an inside of said unit;
decompress-releasing-means for restoring a decompressed condition in said unit to atmos- 50
pheric pressure;
- a bodily-fluids-oozing-device;
a specimen collector disposed in said unit and facing the opening;
a sensor, having an electrode system for contacting with a specimen supplied from said specimen collector, for outputting information regarding the specimen as an electric signal from the electrode system;
determining means for determining a measuring value of the specimen based on the electric signal;
a controller for controlling said humidifying means, said decompressing means, said decompress-releasing-means and said sensor, wherein said system allows said bodily-fluids-oozing-device to function after said humidifying means operates, and also allows said decompressing means to function after said bodily-fluids-oozing-device operates as well as said decompress-releasing-means to function after said sensor outputs the electric signal. 55

FIG. 1

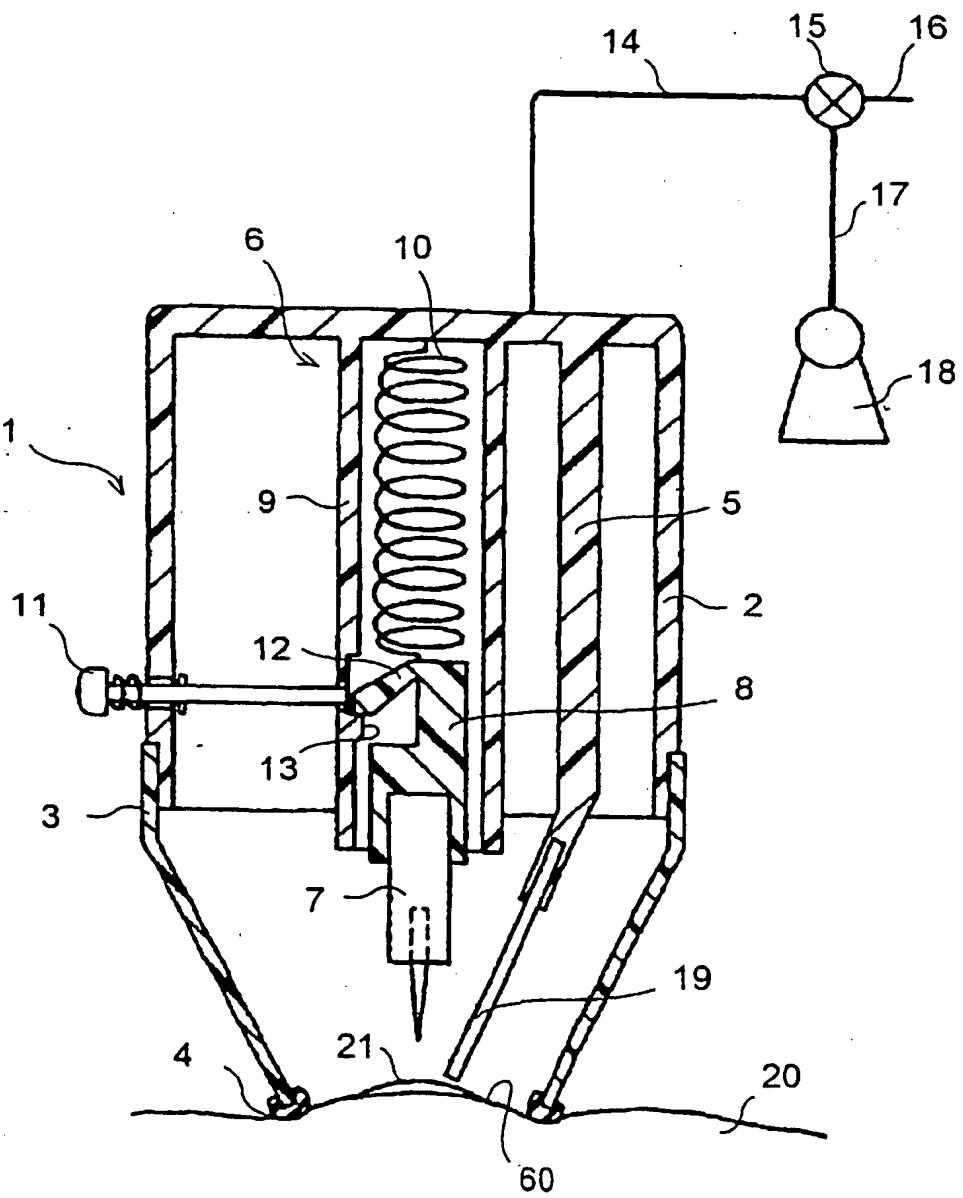


FIG. 2

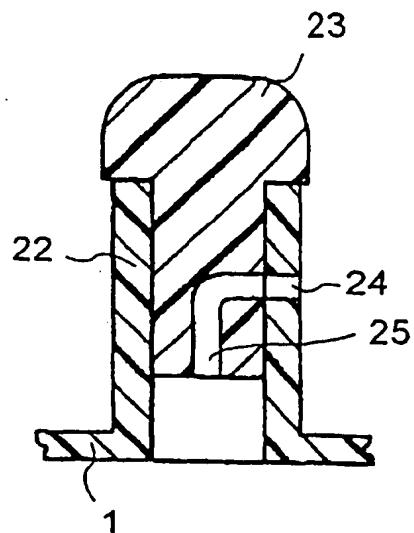


FIG. 3

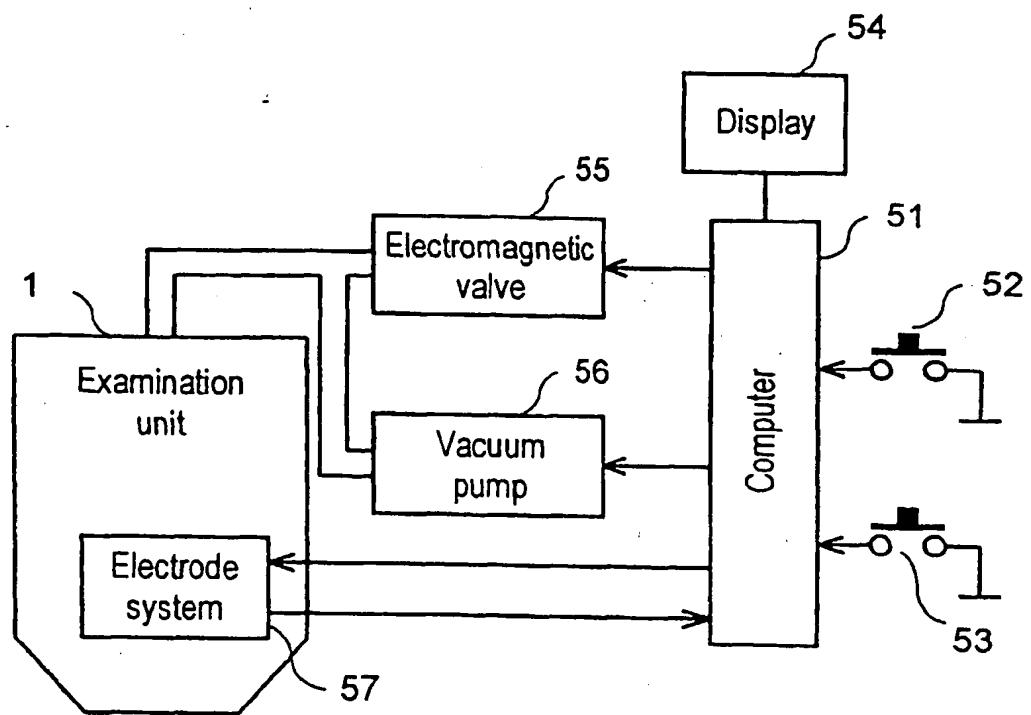


FIG. 4

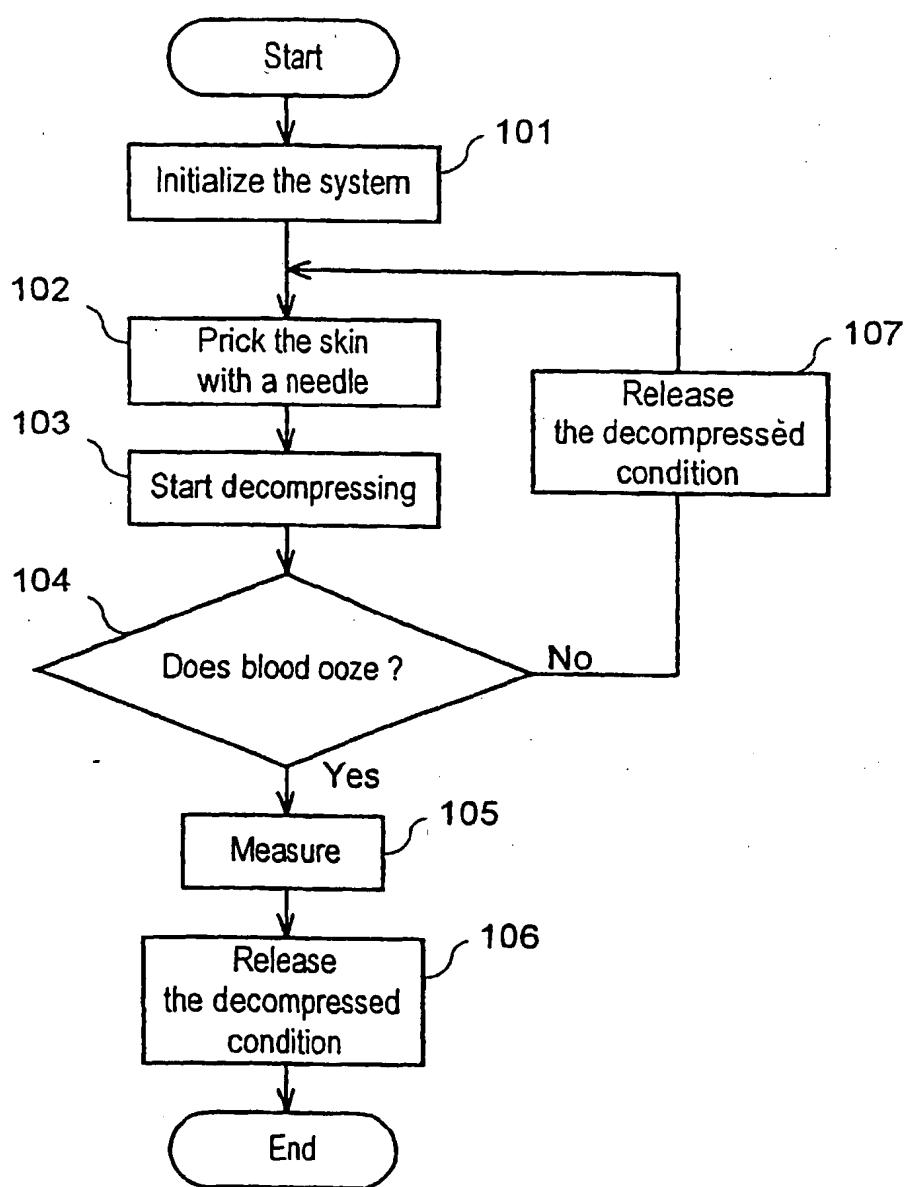


FIG. 5

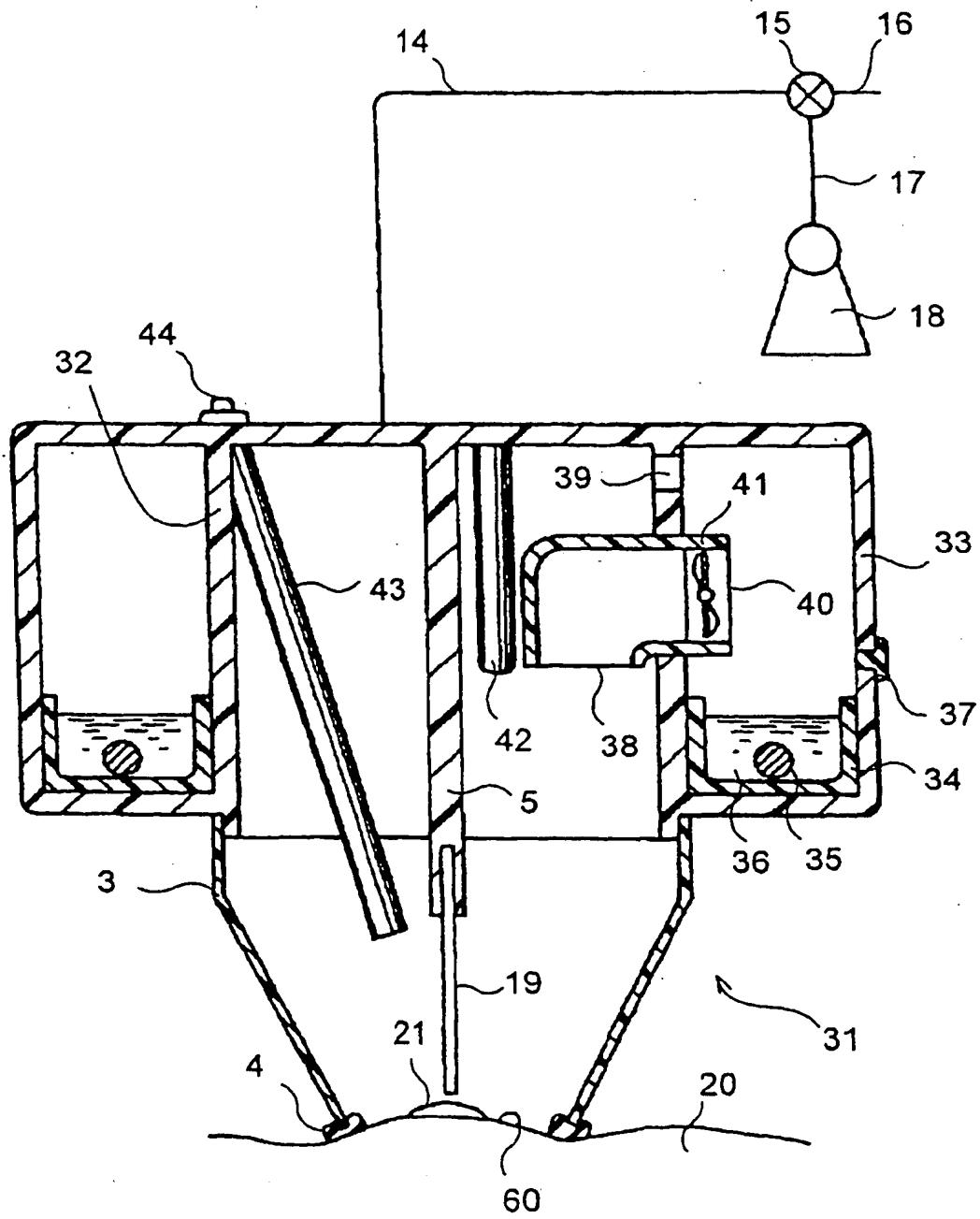


FIG. 6

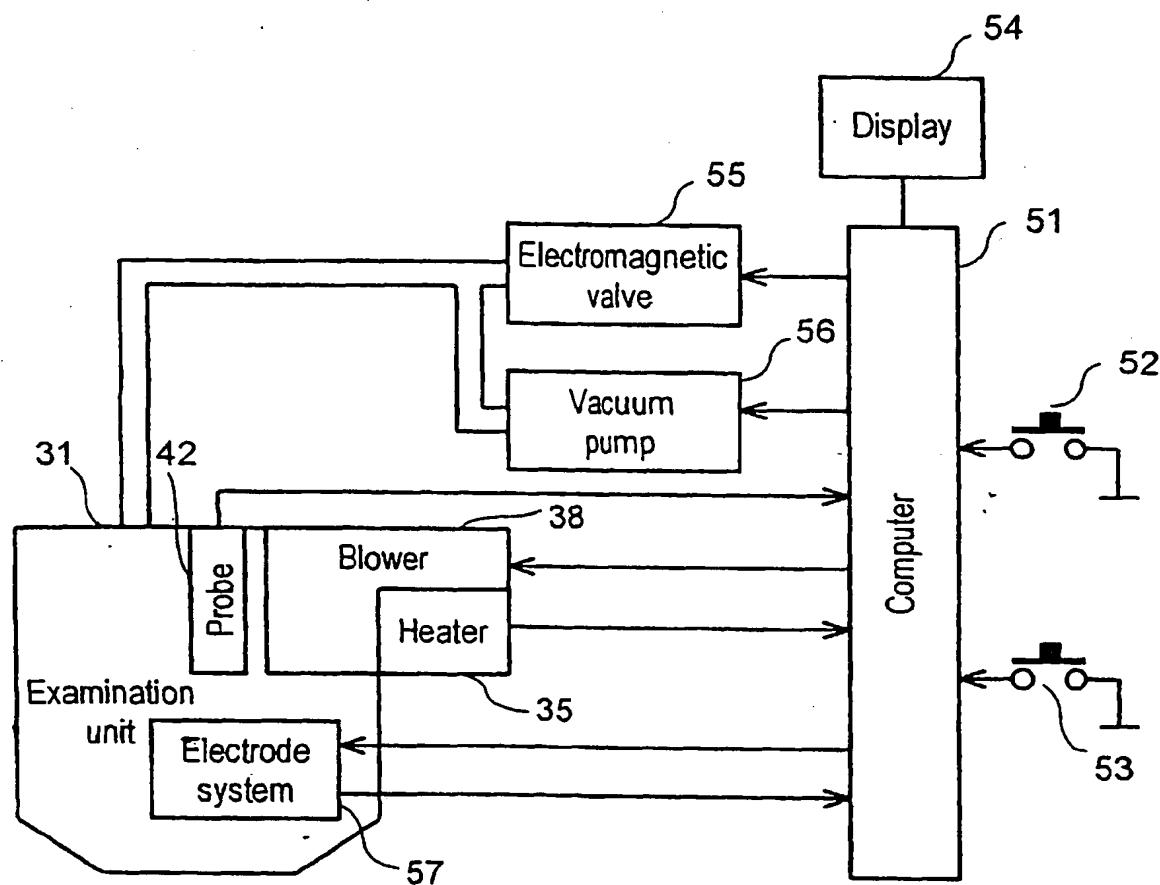
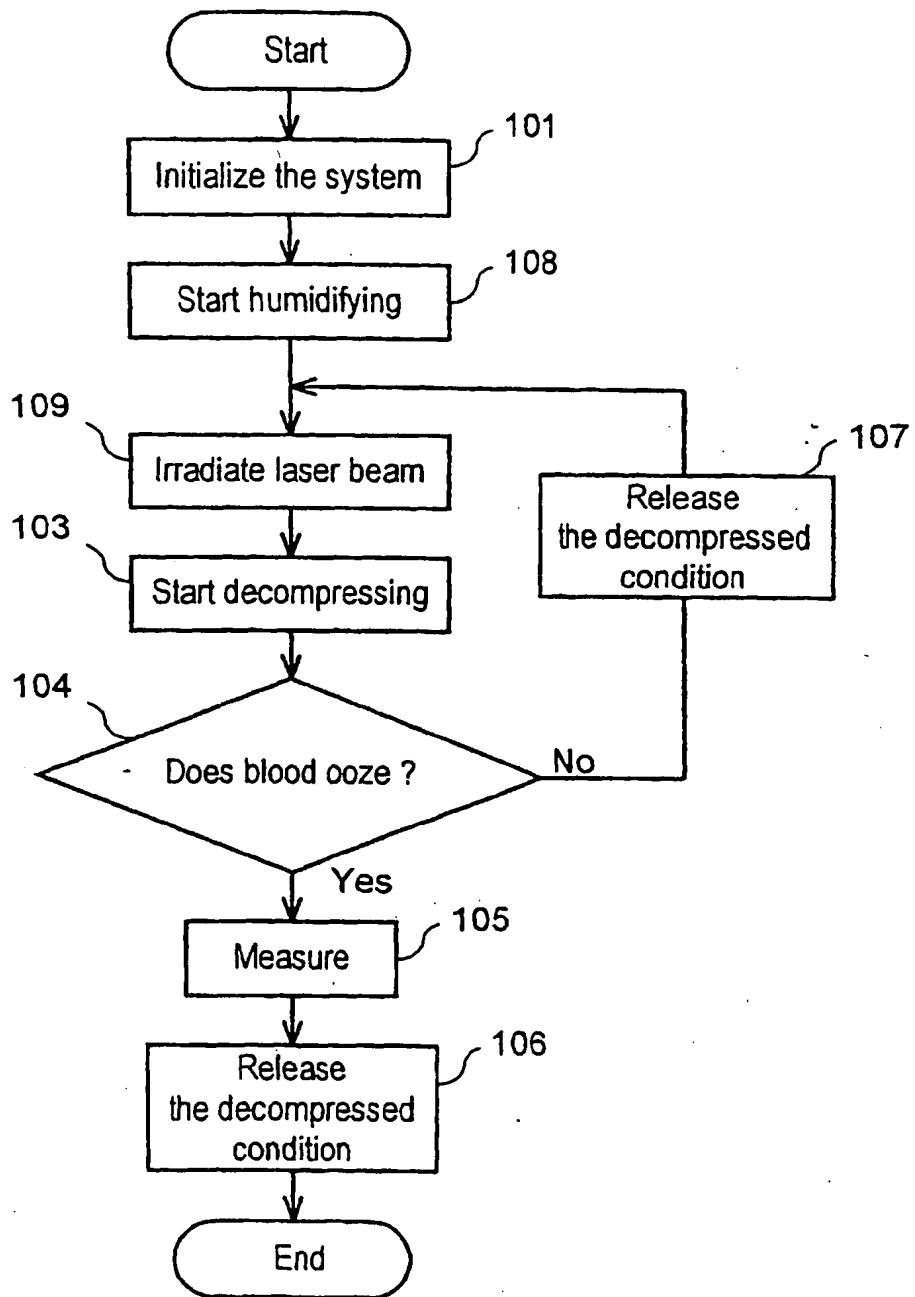


FIG. 7



INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP99/01723

A. CLASSIFICATION OF SUBJECT MATTER
Int.C1⁶ A61B5/14

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
Int.C1⁶ A61B5/14, 10/00Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched -
Jitsuyo-Shinan Koho 1922-1996 Toroku-Jitsuyo-Shinan Koho 1994-1999
Kokai Jitsuyo-Shinan Koho 1971-1999 Jitsuyo-Shinan-Toroku Koho 1996-1999

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages ..	Relevant to claim No.
X	JP, 2-95352, A (K.K. Advance), 6 April, 1990 (06. 04. 90), Full text ; Figs. 1, 2	1-2, 6
Y	Full text ; Figs. 1, 2	5, 7
A	Full text ; Figs. 1, 2 (Family: none)	3-4
Y	Microfilm of the specification and drawings annexed to the request of Japanese Utility Model Application No. 1-49596. (Laid-open No. 2-141414). (NEC Corp.), 28 November, 1990 (28. 11. 90), Page 12, line 17 to page 13, line 10 ; Figs. 3, 4	5, 7
A	Page 12, line 17 to page 13, line 10 ; Figs. 3, 4 (Family: none)	8-10
A	JP, 9-285459, A (Casio Computer Co., Ltd.), 4 November, 1997 (04. 11. 97), Claims ; Par. No. [0027] ; Figs. 1, 2 (Family: none)	8-10

 Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A"	"X"	earlier document but published on or after the international filing date
"E"	"Y"	document which may throw doubt on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
"L"	"&"	document referring to an oral disclosure, use, exhibition or other means
"O"		document published prior to the international filing date but later than the priority date claimed
"P"		

Date of the actual completion of the international search 7 June, 1999 (07. 06. 99)	Date of mailing of the international search report 15 June, 1999 (15. 06. 99)
Name and mailing address of the ISA/ Japanese Patent Office	Authorized officer
Faximile No.	Telephone No.

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